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Notice of Allowability	Application No.	Applicant(s)
	10/088,724	NISHIMOTO, IKUO
	Examiner	Art Unit
	Olga N. Chernyshev	1649
The MAILING DATE of this communication app All claims being allowable, PROSECUTION ON THE MERITS Is herewith (or previously mailed), a Notice of Allowance (PTOL-8: NOTICE OF ALLOWABILITY IS NOT A GRANT OF PATENT of the Office or upon petition by the applicant. See 37 CFR 1.3	S (OR REMAINS) CLOSED in 5) or other appropriate commur RIGHTS. This application is su	this application. If not included nication will be mailed in due course. THIS
1. This communication is responsive to <u>July 13, 2007</u> .		
2. The allowed claim(s) is/are <u>1,2,4-8,13,20-22,27-30,35-38</u>	3 <u>,43 and 45</u> .	
 3.	ve been received. ve been received in Application	ı No
* Certified copies not received:		
Applicant has THREE MONTHS FROM THE "MAILING DATE noted below. Failure to timely comply will result in ABANDON THIS THREE-MONTH PERIOD IS NOT EXTENDABLE.	of this communication to file a MENT of this application.	a reply complying with the requirements
4. A SUBSTITUTE OATH OR DECLARATION must be sub- INFORMAL PATENT APPLICATION (PTO-152) which gi	mitted. Note the attached EXAI ves reason(s) why the oath or	MINER'S AMENDMENT or NOTICE OF declaration is deficient.
5. CORRECTED DRAWINGS (as "replacement sheets") me	ust be submitted.	
(a) ☐ including changes required by the Notice of Draftspe		(PTO-948) attached
1) 🔲 hereto or 2) 🔲 to Paper No./Mail Date		
(b) ☐ including changes required by the attached Examine Paper No./Mail Date	r's Amendment / Comment or i	n the Office action of
Identifying indicia such as the application number (see 37 CFR each sheet. Replacement sheet(s) should be labeled as such in	1.84(c)) should be written on the the header according to 37 CFR	e drawings in the front (not the back) of t 1.121(d).
6. DEPOSIT OF and/or INFORMATION about the dep attached Examiner's comment regarding REQUIREMENT	osit of BIOLOGICAL MATE TFOR THE DEPOSIT OF BIO	RIAL must be submitted. Note the LOGICAL MATERIAL.
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Attachment(s)		
1. Notice of References Cited (PTO-892)		ormal Patent Application
2. Notice of Draftperson's Patent Drawing Review (PTO-948)		mmary (PTO-413), fail Date
☐ Information Disclosure Statements (PTO/SB/08), Paper No./Mail Date	7. 🔲 Examiner's A	mendment/Comment
Examiner's Comment Regarding Requirement for Deposit of Biological Material	8. 🗌 Examiner's S	statement of Reasons for Allowance
	9. 🔲 Other	OLGA M. CHERAYSHEVPH D
		PRIMARY EXAMINER

Amendments To The Claims:

This listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

1. (Previously presented) An isolated polypeptide that suppresses neuronal death associated with Alzheimer's disease having an amino acid sequence of Formula (I):

Pro-Xn₁-(Cys/bXaa)-(Leu/Arg)-Xn₂-Leu-Thr-(Gly/Ser)-Xn₃-Pro (I) (SEQ ID NO: 63)

wherein "Cys/bXaa" indicates Cys or a basic amino acid; "(Leu/Arg)" indicates Leu or Arg; "(Gly/Ser)" indicates Gly or Ser; and Xn₁, Xn₂, and Xn₃ independently indicate arbitrary amino acid sequences not more than 10 residues in length, respectively.

- 2. (Previously presented) An isolated polypeptide selected from the group consisting of:
- (a) a polypeptide having an amino acid sequence selected from the group consisting of SEQ ID NOs: 5 to 8, 10, 12, 13, 21 to 24, 26 to 29, 32, 33, 37 to 40, 46, 48, 54, and 60 and
- (b) a polypeptide that suppresses neuronal death associated with Alzheimer's disease having an amino acid sequence which differs from a polypeptide of SEQ ID NOs: 5 to 8, 10, 12, 13, 21 to 24, 26 to 29, 32, 33, 37 to 40, 46, 48, 54, and 60, in such a way that one amino acid has been substituted, deleted, inserted, or added.

3. (Canceled)

3 M. (Previously presented) A fusion polypeptide comprising the polypeptide of any of claims 1 to 2 fused with one or more other polypeptides.

- (Previously presented) An isolated DNA encoding a polypeptide selected from the group consisting of:
- (a) a polypeptide that suppresses neuronal death associated with Alzheimer's disease having the amino acid sequence of Formula (I):

Pro-Xn₁-(Cys/bXaa)-(Leu/Arg)-Xn₂-Leu-Thr-(Gly/Ser)-Xn₃-Pro (I) (SEQ ID NO: 63)

wherein "Cys/bXaa" indicates Cys or a basic amino acid; "(Leu/Arg)" indicates Leu or Arg; "(Gly/Ser)" indicates Gly or Ser; and Xn₁, Xn₂, and Xn₃ independently indicate arbitrary amino acid sequences not more than 10 residues in length, respectively;

- (b) a polypeptide comprising an amino acid sequence which differs from a polypeptide of SEQ ID NOs: 5 to 8, 10, 12, 13, 21 to 24, 26 to 29, 32, 33, 37 to 40, 46, 48, 54, and 60 in such a way that one amino acid has been substituted, deleted, inserted, or added, wherein the polypeptide suppresses neuronal death associated with Alzheimer's disease;
- (c) a polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 5 to 8, 10, 12, 13, 21 to 24, 26 to 29, 32, 33, 37 to 40, 46, 48, 54, and 60; and
- (d) a fusion polypeptide comprising the polypeptide of (a) or (c) fused with one or more other polypeptides;

wherein the DNA does not comprise the sequence of SEQ ID NO:4.

- 6 (Previously presented) A vector into which a DNA encoding a polypeptide of any one of (a) to (c) is inserted:
- (a) a polypeptide that suppresses neuronal death associated with Alzheimer's disease having the amino acid sequence of Formula (I):

Pro-Xn₁-(Cys/bXaa)-(Leu/Arg)-Xn₂-Leu-Thr-(Gly/Ser)-Xn₃-Pro (I) (SEQ ID NO: 63)

wherein "Cys/bXaa" indicates Cys or a basic amino acid; "(Leu/Arg)" indicates Leu or Arg; "(Gly/Ser)" indicates Gly or Ser; and Xn₁, Xn₂, and Xn₃ independently indicate arbitrary amino acid sequences not more than 10 residues in length, respectively;

- (b) a polypeptide comprising an amino acid sequence which differs from a polypeptide of SEQ ID NOs: 5 to 8, 10, 12, 13, 21 to 24, 26 to 29, 32, 33, 37 to 40, 46, 48, 54, and 60 in such a way that one amino acid has been substituted, deleted, inserted, or added, wherein the polypeptide suppresses neuronal death associated with Alzheimer's disease;
- (c) a polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 5 to 8, 10, 12, 13, 21 to 24, 26 to 29, 32, 33, 37 to 40, 46, 48, 54, and 60; and
- (d) a fusion polypeptide comprising the polypeptide of (a) or (b) fused with one or more other polypeptides.
 - 6 A. (Original) A host cell retaining the vector of claim 8.
- (Previously presented) A method for producing the polypeptide of any one of claims 1 to 2 or a fusion polypeptide comprising the polypeptide of any one of claims 1 to 2, comprising:

culturing a host cell retaining a vector into which a DNA encoding the polypeptide of any one of claims 1 to 2, or a fusion polypeptide comprising the polypeptide of any one of claims 1 to 2 fused with one or more other polypeptides, is inserted; and recovering an expressed polypeptide from the host cell or culture supernatant thereof.

9-12. (Canceled)

(Previously presented) A pharmaceutical composition comprising the polypeptide of any one of claims 1 to 2.

14-16. (Canceled)

17-19. (Canceled)

- Q 20. (Previously presented) The polypeptide of claim 1, wherein Xn_1 is an amino acid sequence consisting of 3 to 5 arbitrary amino acids, Xn_2 is an amino acid sequence consisting of 1 to 3 arbitrary amino acids, and Xn_3 is an amino acid sequence consisting of 3 to 5 arbitrary amino acids.
- (Previously presented) The polypeptide of claim 1, wherein the polypeptide comprises an amino acid sequence of SEQ ID NO: 101.
- [Note that the polypeptide of claim 1, wherein the polypeptide comprises an amino acid sequence of SEQ ID NO: 102.

23-26. (Canceled)

- (Previously presented) The polypeptide of claim 2, wherein the polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 5 to 8, 10, 12, 13, 21 to 24, 26 to 29, 32, 33, 37 to 40, 46, 48, 54, and 60.
- 13 28. (Previously presented) The DNA of claim 5, wherein Xn₁ is an amino acid sequence consisting of 3 to 5 arbitrary amino acids, Xn₂ is an amino acid sequence consisting of 1 to 3 arbitrary amino acids, and Xn₃ is an amino acid sequence consisting of 3 to 5 arbitrary amino acids.
- (Previously presented) The DNA of claim 5, wherein the DNA encodes a polypeptide comprising an amino acid sequence of SEQ ID NO: 101.
- 15 30. (Previously presented) The DNA of claim 5, wherein the DNA encodes a polypeptide comprising an amino acid sequence of SEQ ID NO: 102.

31-34. (Canceled)

- (Previously presented) The DNA of claim 5, wherein the DNA encodes a polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 6 to 8, 10, 24, 26 to 29, 32, 33, 37 to 40, 46, 48, 54, and 60.
- (Previously presented) The vector of claim 6, wherein Xn₁ is an amino acid sequence consisting of 3 to 5 arbitrary amino acids, Xn₂ is an amino acid sequence consisting of 1 to 3 arbitrary amino acids, and Xn₃ is an amino acid sequence consisting of 3 to 5 arbitrary amino acids.
- (Previously presented) The vector of claim 6, wherein the DNA encodes a polypeptide comprising an amino acid sequence of SEQ ID NO: 101.
- 19 38. (Previously presented) The vector of claim 6, wherein the DNA encodes a polypeptide comprising an amino acid sequence of SEQ ID NO: 102.

39-42. (Canceled)

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- QD A3. (Previously presented) The vector of claim 6, wherein the DNA encodes a polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 5 to 8, 10, 12, 13, 21 to 24, 26 to 29, 32, 33, 37 to 40, 46, 48, 54, and 60.
 - 44. (Canceled)
- 2\ A5. (Previously presented) A composition comprising a polypeptide of claim 2, and a carrier.

46-49. (Canceled)